IN THE UNITED STATES DISTRICT COURT FOR THE SOUTHERN DISTRICT OF NEW YORK

BERTHA S. FARMER,

Plaintiff,

Civil Action No.: 1:08-00-3058

MERCK & CO., INC.,

Defendant.

COMPLAINT AND DEMAND FOR JURY TRIAL

Plaintiff, BERTHA S. FARMER, through her undersigned attorneys Levin, Papantonio et al., sues Defendant Merck & Company, Inc., and alleges as follows:

I. JURISDICTION AND VENUE

- This Court has jurisdiction pursuant to 28 U.S.C. §§1332, as complete diversity 1. exists between Plaintiff and Defendant. Plaintiff is a resident of the State of Georgia, and Defendant is incorporated and has its primary place of business in the State of New Jersey. The amount in controversy, exclusive of interest and costs, exceeds \$75,000.
- Venue is proper within this district pursuant to Case Management Order No. 3, filed 2. November 1, 2006, signed by John F. Keenan, allowing Fosamax-related cases to be filed directly in the Southern District of New York.

II. PARTIES

- Plaintiff BERTHA S. FARMER was born October 17, 1928. At all relevant times 3. Plaintiff was a resident of the State of Georgia, and used FOSAMAX from approximately August 2003 through at least May 2006.
- Defendant is a corporation organized and existing under the laws of the State of New 4. Jersey, with its principal place of business in New Jersey. The Defendant's registered office is at 820 Bear Tavern Road, City of West Trenton, Mercer County, New Jersey.
- Defendant was at all relevant times authorized to conduct business in the State of 5. Georgia.
- At all times relevant Defendant regularly transacted business in the State of Georgia 6. and continues to do so.
- At all relevant times Defendant, through its agents, servants, employees and apparent 7. agents was the designer, manufacturer, marketer, distributor and seller of FOSAMAX, a bisphosphonate drug used primarily to mitigate or reverse the effects of osteoporosis.
- Defendant, either directly or through its agents, apparent agents, servants or 8. employees, at all relevant times, sold and distributed FOSAMAX in the State of Georgia for the treatment or prevention of osteoporosis, Paget's Disease and other off-label uses.
- Defendant derives substantial revenue from pharmaceutical products used or 9.

- consumed in the State of Georgia.
- 10. Defendant expected, or should have expected, that its business activities could or would have consequences within the State of Georgia.

III. SUMMARY OF THE CASE

- Defendant, either directly or through its agents, apparent agents, servants or employees designed, manufactured, marketed, advertised, distributed and sold FOSAMAX for the treatment of osteoporosis, Paget's Disease, and other off-label uses.
- 12. As a result of the defective nature of FOSAMAX, persons who were prescribed and ingested FOSAMAX, including Plaintiff BERTHA S. FARMER, have suffered and may continue to suffer severe and permanent personal injuries to the jaw bone, including osteonecrosis of the jaw and other diagnoses of irreversible damage to the jaw.
- 13. Defendant concealed its knowledge of FOSAMAX's unreasonably dangerous risks from Plaintiff BERTHAS. FARMER, other consumers, and the medical community.
- 14. Defendant failed to conduct adequate and sufficient post-marketing surveillance of FOSAMAX after it began marketing, advertising, distributing, and selling the drug.
- 15. As a result of Defendant's actions and inaction, Plaintiff BERTHA S. FARMER was injured due to her ingestion of FOSAMAX, which has caused and will continue to cause Plaintiff's various injuries and damages. Plaintiff accordingly seeks

compensatory damages.

IV. FACTUAL BACKGROUND

- 16. At all relevant times Defendant was responsible for, or involved in, designing, manufacturing, marketing, advertising, distributing, and selling FOSAMAX.
- 17. In September 1995, the United States Food and Drug Administration ("FDA") approved Merck's compound alendronate, which is marketed by Merck as FOSAMAX, for various uses, including the treatment of osteoporosis and Paget's Disease.
- 18. FOSAMAX falls within a class of drugs known as bisphosphonates.

 Bisphosphonates are used for treating bone conditions such as osteoporosis and Paget's disease. Other drugs within this class such as Aredia and Zometa are also used as chemotherapy and as adjunct chemotherapy but are not indicated for use in non-cancerous conditions such as osteoporosis.
- 19. There are two classes of bisphosphonates: the N-containing (nitrogenous) and non-N-containing (non-nitrogenous) bisphosphonates. The nitrogenous bisphophonates include the following: pamidronate (Aredia); ibandronate (Boniva); risedronate (Actonel); and alendronate (FOSAMAX). The non-nitrogenous bisphosphonates include the following: etridonate (Didronel); clodronate (Bonefos and Loron); and tiludronate (Skelid). Alendronate, like the others, contains a nitrogen atom, whereas etridonate, clodronate, and tiludronate do not. The PDR for FOSAMAX confirms

that the molecule contains a nitrogen atom.

- 20. Throughout the 1990s and 2000s, medical articles and studies appeared reporting the frequent and common occurrence of osteonecrosis of the jaw within the nitrogenous bisphosphonates used for chemotherapy. As with its reported and acknowledged side effects concerning irritation, erosion, and inflammation of the upper gastrointestinal tract, Merck knew or should have know that FOSAMAX, as a nitrogenous bisphosphonate, shared a similar adverse event profiles to the other drugs within this specific subclass of bisphosphonates (i.e., those containing nitrogen).
- Merck knew and or should have known that bisphosphonates, including FOSAMAX, 21. inhibit endothelial cell function. Similarly, Merck knew or should have known that bisphosponates also inhibit vascularization of the affected area and induce ischemic changes specific to patients mandibles (lower jaws) and maxillae (upper jaws) and that these ischemic changes appear to be cumulative in nature.
- 22. Merck also knew or should have known that these factors combine to create a compromised vascular supply in the affected area. As a result, a minor injury or disease can turning into a non-healing wound. That in turn can progress to widespread necrosis (bone death) and osteomyelitis (inflammation of bone marrow).
- Dentists are now being advised by state dental associations to refrain from using any 23. invasive procedure (such as drilling a cavity) for any patient on FOSAMAX.
- Once the osteonecrosis begins and becomes symptomatic, it is very difficult to treat 24. and is not reversible.

- 25. Shortly after Defendant began selling FOSAMAX, reports of osteonecrosis of the jaw and other dental complications among users began surfacing, indicating that FOSAMAX shared the class effects of the other nitrogenous bisphosphonates. Despite this knowledge, Defendant failed to implement further study risk of osteonecrosis of the jaw relative to FOSAMAX. Rather than evaluating and verifying the safety of FOSAMAX with respect to osteonecrosis of the jaw, Defendant proposed further uses of FOSAMAX, such as FOSAMAX-D, and sought to extend the exclusivity period of FOSAMAX through 2018.
- 26. Osteonecrosis of the jaw is a serious medical event and can result in severe disability and death.
- 27. Since FOSAMAX was released, the FDA has received a number of reports osteonecrosis of the jaw among users of FOSAMAX.
- 28. On August 25, 2004, the FDA posted its ODS (Office of Drug Safety) Postmarketing Safety Review on bisphosphonates - specifically pamidronate (Aredia), zoledronic acid (Zometa), risedronate (Actonel), and alendronate (FOSAMAX). This was an epidemiologic review of the FDA adverse events database conducted by the FDA's Division of Drug Risk Evaluation.
- 29. As a result of the FDA Review, the FDA observed that the risk of osteonecrosis of the jaw was not confined to bisphosphonates used for chemotherapy. The FDA's review indicated that the osteonecrosis of the jaw was a class effect which specifically extended to the oral bisphosphonate, FOSAMAX.

- 30. As a result, the FDA recommended and stated that the labeling for FOSAMAX should be amended by Merck to specifically warn about the risk of osteonecrosis of the jaw. Merck has refused to accede to the FDA's request and, to this day, still does not warn of the risk of osteonecrosis of the jaw in its FOSAMAX labeling.
- 31. Rather than warn patients, and despite knowledge known by Defendant about increased risk of osteonecrosis of the jaw in patients using FOSAMAX, Defendant continues to defend FOSAMAX and minimize unfavorable findings.
- 32. FOSAMAX is one of Defendant's top selling drugs. Averaging more than \$3 billion a year in sales.
- 33. Consumers, including Plaintiff BERTHA S. FARMER, who have used FOSAMAX for the treatment or prevention of osteoporosis, Paget's Disease and/or other off-label uses, have several alternative safer products available to treat their conditions.
- Defendant knew of the significant risk of dental and oral complications caused by ingestion of FOSAMAX, but Defendant did not adequately and sufficiently warn consumers, including Plaintiff BERTHA S. FARMER, or the medical community, of such risks.
- As a direct result, Plaintiff BERTHA S. FARMER was prescribed FOSAMAX and has been permanently and severely injured, having suffered serious consequences from the ingestion of FOSAMAX. Plaintiff BERTHA S. FARMER requires and will in the future require ongoing medical care and treatment.
- 36. Plaintiff BERTHA S. FARMER has suffered from mental anguish from the

- knowledge that she will have life-long complications as a result of the injuries Plaintiff sustained from the use of FOSAMAX.
- 37. Plaintiff BERTHA S. FARMER was prescribed and began taking FOSAMAX in August 2003.
- 38. Plaintiff used FOSAMAX as prescribed and in a foreseeable manner.
- 39. As a direct and proximate result of using FOSAMAX, Plaintiff suffered severe personal injury to her jaw.
- 40. Plaintiff, as a direct and proximate result of using FOSAMAX, suffered severe mental and physical pain and suffering and has sustained permanent injuries and emotional distress.
- 41. Plaintiff used FOSAMAX which had been provided to her in a condition that was substantially the same as the condition in which it was manufactured and sold.
- 42. Plaintiff would not have used FOSAMAX had Defendant properly disclosed the risks associated with the drug. Alternatively, Plaintiff would have known the precursor events of osteonecrosis of the jaw and would have been able to avoid the clinical manifestation of the symptoms as they currently exist.
- 43. Defendant, through its affirmative misrepresentations and omissions, actively concealed from Plaintiff and her physicians the true and significant risks associated with taking FOSAMAX. The running of any applicable statute of limitations has been tolled by reason of Defendant's fraudulent concealment.
- 44. As a result of Defendant's actions, Plaintiff and her prescribing physicians were

unaware, and could not have reasonably known or have learned through reasonable diligence, that Plaintiff had been exposed to the risks identified in this complaint, and that those risks were the direct and proximate result of Defendant's acts, omissions, and misrepresentations.

V. COUNTS

COUNT I: NEGLIGENCE

- 45. Plaintiff re-alleges the above paragraphs as if fully set forth herein.
- 46. Defendant owed Plaintiff, BERTHA S. FARMER, and other consumers, a duty to exercise reasonable care when designing, manufacturing, marketing, advertising, distributing, and selling FOSAMAX.
- 47. Defendant failed to exercise due care under the circumstances and therefore breached this duty by:
- a. failing to properly and thoroughly test FOSAMAX before releasing the drug to market;
- b. failing to properly and throughly analyze the data resulting from the pre-marketing tests of FOSAMAX;
 - c. failing to conduct sufficient post-market testing and surveillance of FOSAMAX;
- d. designing, manufacturing, marketing, advertising, distributing, and selling FOSAMAX to consumers, including Plaintiff, without an adequate warning of the significant and dangerous risks of FOSAMAX and without proper instructions to avoid the harm which could foreseeably occur as a result of using the drug;

- e. failing to exercise due care when advertising and promoting FOSAMAX; and
- f. negligently continuing to manufacture, market, advertise, and distribute FOSAMAX after Defendant knew or should have known of its adverse effects.
- As a direct and proximate consequence of Defendant's actions, omissions, and misrepresentations, Plaintiff BERTHA S. FARMER sustained significant and permanent injury to her jaw. In addition, Plaintiff required and will continue to require healthcare and services. Plaintiff has incurred and will continue to incur medical and related expenses. Plaintiff also has suffered and will continue to suffer diminished capacity for the enjoyment of life, a diminished quality of life, increased risk of premature death, aggravation of preexisting conditions and activation of latent conditions, and other losses and damages. Plaintiff's direct medical losses and costs include care for hospitalization, physician care, monitoring, treatment, medications, and supplies. Plaintiff has incurred and will continue to incur mental and physical pain and suffering and loss of wages and wage-earning capacity.
- 49. Defendant's conduct as described above was committed with knowing, conscious, wanton, willful, and deliberate disregard for the value of human life and the rights and safety of consumers such as Plaintiff, thereby entitling Plaintiff to punitive damages so as to punish Defendant and deter it from similar conduct in the future.

COUNT II: STRICT LIABILITY

- 50. Plaintiff re-alleges the above paragraphs as if fully set forth herein.
- 51. Defendant manufactured, sold, distributed, marketed, and/or supplied FOSAMAX

- in a defective and unreasonably dangerous condition to consumers, including Plaintiff BERTHA S. FARMER.
- 52. Defendant designed, manufactured, sold, distributed, supplied, marketed, and/or promoted FOSAMAX, which was expected to reach and did in fact reach consumers, including Plaintiff, without substantial change in the condition in which it was manufactured and sold by Defendant.
- 53. Plaintiff used FOSAMAX as prescribed and in a manner normally intended, recommended, promoted, and marketed by Defendant.
- 54. FOSAMAX failed to perform safely when used by ordinary consumers, including Plaintiff, including when it was used as intended and in a reasonably foreseeable manner.
- 55. FOSAMAX was defective in its design and was unreasonably dangerous in that its unforeseeable risks exceeded the benefits associated with its design or formulation.
- 56. FOSAMAX was defective in design or formulation in that it posed a greater likelihood of injury than other similar medications and was more dangerous than an ordinary consumer could reasonably foresee or anticipate.
- 57. FOSAMAX was defective in its design and was unreasonably dangerous in that it neither bore nor was packaged with nor accompanied by warnings adequate to alert consumers, including Plaintiff, of the risks described herein, including, but not limited to, the risk of osteonecrosis of the jaw.
- 58. Although Defendant knew or should have known of the defective nature of

FOSAMAX, it continued to design, manufacture, market, and sell FOSAMAX so as to maximize sales and profits at the expense of the public health and safety. By so acting, Defendant acted with conscious and deliberate disregard of the foreseeable harm caused by FOSAMAX.

- 59. Plaintiff could not, through the exercise of reasonable care, have discovered FOSAMAX's defects or perceived the dangers posed by the drug.
- As a direct and proximate consequence of Defendant's conduct, Plaintiff BERTHA

 S. FARMER sustained significant and permanent injury to her jaw. In addition,
 Plaintiff required and will continue to require healthcare as a result of her injury.

 Plaintiff has incurred and will continue to incur medical and related expenses.

 Plaintiff also has suffered and will continue to suffer diminished capacity for the enjoyment of life, a diminished quality of life, increased risk of premature death, aggravation of preexisting conditions and activation of latent conditions, and other losses and damages. Plaintiff's direct medical losses and costs include care for hospitalization, physician care, monitoring, treatment, medications, and supplies.

 Plaintiff has incurred and will continue to incur mental and physical pain and suffering and loss of wages and wage-earning capacity.
- Defendant's conduct as described above was committed with knowing, conscious, wanton, willful, and deliberate disregard for the value of human life and the rights and safety of consumers such as Plaintiff, thereby entitling Plaintiff to punitive damages so as to punish Defendant and deter it from similar conduct in the future.

COUNT III: BREACH OF EXPRESS WARRANTY

- 62. Plaintiff re-alleges the above paragraphs as if fully set forth herein.
- 63. Defendant expressly represented to Plaintiff BERTHAS. FARMER, other consumers and the medical community that FOSAMAX was safe and fit for its intended purposes, was of merchantable quality, did not produce any dangerous side effects. and had been adequately tested.
- FOSAMAX does not conform to Defendant's express representations because it is 64. not safe, has numerous and serious side effects, and causes severe and permanent injuries.
- At all relevant times FOSAMAX did not perform as safely as an ordinary consumer 65. would expect, when used as intended or in a reasonably foreseeable manner.
- 66. Plaintiff BERTHAS, FARMER, other consumers, and the medical community relied upon Defendant's express warranties.
- As a direct and proximate result of Defendant's actions, Plaintiff BERTHA S. 67. FARMER sustained serious significant and permanent injury to her jaw. In addition, Plaintiff required and will continue to require healthcare and services as a result of her injury. Plaintiff has incurred and will continue to incur medical and related expenses as a result of her injury. Plaintiff also has suffered and will continue to suffer diminished capacity for the enjoyment of life, a diminished quality of life, increased risk of premature death, aggravation of preexisting conditions and activation of latent conditions, and other losses and damages. Plaintiff's direct

medical losses and costs include care for hospitalization, physician care, monitoring, treatment, medications, and supplies. Plaintiff has incurred and will continue to incur mental and physical pain and suffering and loss of wages and wage-earning capacity.

Defendant's conduct as described above was committed with knowing, conscious, 68. wanton, willful, and deliberate disregard for the value of human life and the rights and safety of consumers such as Plaintiff, thereby entitling Plaintiff to punitive damages so as to punish Defendant and deter it from similar conduct in the future.

COUNT IV: BREACH OF IMPLIED WARRANTY

- 69. Plaintiff re-alleges the above paragraphs as if fully set forth herein.
- 70. Defendant manufactured, distributed, advertised, promoted, and sold FOSAMAX.
- At all relevant times, Defendant knew of the use for which FOSAMAX was intended 71. and impliedly warranted the product to be of merchantable quality and safe and fit for such use.
- 72. Defendant was aware that consumers, including Plaintiff BERTHA S. FARMER, would use FOSAMAX for treatment or prevention of osteoporosis or Paget's Disease and for other off-label purposes.
- 73. Plaintiff and the medical community reasonably relied upon the judgment and sensibility of Merck to sell FOSAMAX only if it was indeed of merchantable quality and safe and fit for its intended use.
- 74. Defendant breached its implied warranty to consumers, including Plaintiff; FOSAMAX was not of merchantable quality or safe and fit for its intended use.

- 75. Consumers, including Plaintiff, and the medical community, reasonably relied upon Defendant's implied warranty for FOSAMAX.
- 76. FOSAMAX reached consumers without substantial change in the condition in which it was manufactured and sold by Defendant.
- 77. As a direct and proximate result of Defendant's action, Plaintiff BERTHA S. FARMER sustained significant and permanent injury to her jaw. In addition, Plaintiff required and will continue to require healthcare and services as a result of her injury. Plaintiff has incurred and will continue to incur medical and related expenses as a result of her injury. Plaintiff also has suffered and will continue to suffer diminished capacity for the enjoyment of life, a diminished quality of life, increased risk of premature death, aggravation of preexisting conditions and activation of latent conditions, and other losses and damages. Plaintiff's direct medical losses and costs include care for hospitalization, physician care, monitoring, treatment, medications, and supplies. Plaintiff has incurred and will continue to incur mental and physical pain and suffering and loss of wages and wage-earning capacity.
- 78. Defendant's conduct as described above was committed with knowing, conscious, wanton, willful, and deliberate disregard for the value of human life and the rights and safety of consumers such as Plaintiff, thereby entitling Plaintiff to punitive damages so as to punish Defendant and deter it from similar conduct in the future.

COUNT V: FRAUDULENT MISREPRESENTATION

79. Plaintiff re-alleges the above paragraphs as if fully set forth herein.

- 80. Defendant made fraudulent misrepresentations with respect to FOSAMAX in the following particulars:
- a.. Defendant represented through its labeling, advertising, marketing materials, detail persons, seminar presentations, publications, notice letters, and regulatory submissions that FOSAMAX had been tested and found to be safe and effective for the treatment of osteoporosis and Paget's Disease; and
- b. Defendant represented that FOSAMAX was safer than other alternative medications.
- Defendant knew that its representations were false, yet it willfully, wantonly, and recklessly disregarded its obligation to provide truthful representations regarding the safety and risk of FOSAMAX to consumers, including Plaintiff, and the medical community.
- 82. The representations were made by Defendant with the intent that doctors and patients, including Plaintiff, rely upon them.
- 83. Defendant's representations were made with the intent of defrauding and deceiving Plaintiff, other consumers, and the medical community to induce and encourage the sale of FOSAMAX.
- 84. Plaintiff BERTHA S. FARMER, Plaintiff's doctors, and others relied upon the representations.
- 85. Defendant's fraudulent representations evinced its callous, reckless, willful, and depraved indifference to the health, safety, and welfare of consumers, including

Plaintiff.

- As a direct and proximate result, Plaintiff BERTHA S. FARMER sustained significant and permanent injury to her jaw. In addition, as a result of her injury, Plaintiff required and will continue to require healthcare and services, and has incurred and will continue to incur medical and related expenses. Plaintiff also suffered and will continue to suffer diminished capacity for the enjoyment of life, a diminished quality of life, increased risk of premature death, aggravation of preexisting conditions and activation of latent conditions, and other losses and damages. Plaintiff's direct medical losses and costs include care for hospitalization, physician care, monitoring, treatment, medications, and supplies. Plaintiff has incurred and will continue to incur mental and physical pain and suffering and loss of wages and wage-earning capacity.
- 87. Defendant's conduct as described above was committed with knowing, conscious, wanton, willful, and deliberate disregard for the value of human life and the rights and safety of consumers such as Plaintiff, thereby entitling Plaintiff to punitive damages so as to punish Defendant and deter it from similar conduct in the future.

COUNT VI: FRAUDULENT CONCEALMENT

- 88. Plaintiff re-alleges the above paragraphs as if fully set forth herein.
- 89. Defendant fraudulently concealed information with respect to FOSAMAX in the following particulars:
 - a. Defendant represented through its labeling, advertising, marketing materials, detail

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- b. Defendant represented that FOSAMAX was safer than other alternative medications and fraudulently concealed information which demonstrated that FOSAMAX was not safer than alternatives available on the market.
- 90. Defendant had sole access to material facts concerning the dangers and unreasonable risks associated with FOSAMAX.
- 91. Defendant's concealment of information about the risks associated with taking FOSAMAX was intentional, and the representations made by Defendant were known by Defendant to be false.
- 92. The concealment of information and the misrepresentations about FOSAMAX were made by Defendant with the intent that doctors and patients, including Plaintiff, rely upon them.
- 93. Plaintiff BERTHA S. FARMER, Plaintiff's doctors, and others relied upon the representations and were unaware of the substantial dental and oral risks associated with taking FOSAMAX that Defendant had concealed from them.
- 94. As a direct and proximate result of Defendant's fraudulent concealment and misrepresentations, Plaintiff BERTHA S. FARMER suffered significant and permanent injury to her jaw as well as severe and permanent injuries, including pain, mental and physical anguish and suffering, a diminished capacity for the enjoyment

of life, aggravation of preexisting conditions and activation of latent conditions, and a fear of developing other harmful conditions or problems as a result of the injury. Plaintiff has suffered and will continue to suffer a loss of wages and wage-earning capacity and has incurred expenses for medical care and treatment due to the injuries caused by FOSAMAX.

95. Defendant's conduct as described above was committed with knowing, conscious, wanton, willful, and deliberate disregard for the value of human life and the rights and safety of consumers such as Plaintiff, thereby entitling Plaintiff to punitive damages so as to punish Defendant and deter it from similar conduct in the future.

GLOBAL PRAYER FOR RELIEF

WHEREFORE, Plaintiff demands judgment against Defendant, as follows:

- a. compensatory damages on each cause of action;
- b. punitive damages on each cause of action;
- c. reasonable attorneys' fees where recoverable;
- d. costs of this action; and
- e. such other additional and further relief as the Court may deem necessary, appropriate, and just.

VI. DEMAND FOR JURY TRIAL

Plaintiff demands a trial by jury on all counts and issues so triable.

MEGHAN M. TANS

TIMOTHY M. O'BRIEN

LEVIN, PAPANTONIO, THOMAS, MITCHELL,

ESCHSNER & PROCTOR, P.A.

316 South Baylen Street, Suite 600 (32502)

P. O. Box 12308

Pensacola, Florida 32591

Telephone(850) 435-7181

FAX (850) 436-6181

JS 44C/SDNY REV. 12/2005

CIVIL COVER SHEET

The JS-44 civil cover sheet and the information contained herein neither replace nor supplement the filing and service of pleadings or other papers as required by law, except as provided by local rules of court. This form, approved by the Judicial Conference of the United States in September 1974, is required for use of the Clerk of Court for the purpose of initiating the civil docket sheet.

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UNITED STATES DISTRICT COURT (NEW YORK SOUTHERN)

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111 EIGHTH AVENUE NEW YORK, NEW YOR	RK, 10011						
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YOU ARE HEREBY SUMMON	ED and requ	uired to serve up	on PLAINTIFF'S	S ATTORNEY (name and address)			
MEGHAN M. TANS LEVIN, PAPANTONIO, 316 S. BAYLEN STREE PENSACOLA, FL 3259	ET, SUITE 60		SNER & PROC	ΓOR, P.A.			
·							
an answer to the complaint which is here summons upon you, exclusive of the dathe relief demanded in the complaint. You of time after service.	ay of service	. If you fail to do	so, judgment b	days after service of this by default will be taken against you for fithis Court within a reasonable period			
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Case '	1:08-cv-03058-JFK
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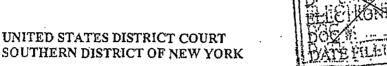
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Case 1:06-md-01789-JFK-JCF

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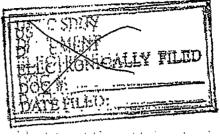
Page 1 of 3



IN RE: Fosamax Products Liability Litigation

This Document Relates to: MDL-1789

Fredericka Allen v. Merck Case No.:06-cv-9452 Bridget Arcemont v. Merck Case No.: 07-cy-2389 Virginia Bergeson v. Merck Case No.: 06-cv-9454 Shirley Boles v. Merck Case No.:06-cy-9455 Sharon Bradford v. Merck Case No.: 07-cv-477 Winnie Bruswell v. Merck Case No.: 1:07-cv-1413 Roberta Brodin v. Merck Case No : 07-cv-3466 Brenda Carpenter v. Merck Case No.: 07-cv-3464 Sonja Daroff v. Merck Case No.: 07-cv-3462 Debra Flores v. Merck Case No.: 07-cv-02442 Sandra Hall v. Merck Case No.: 07-cv-3467 Porcia Hardy v. Merck Case No.: 07-cv-797 Carolyn Hester v. Merck Case No.:06-cy-9450 Puniela Hines v. Merck Case No.: 07-cv-480 Rochelle Kenigav, Merck Case No.: 07-cv-00478 Anna Merwin v. Merck Case No.: 07-cv-3463 Junice Moe v. Merck Case No.: 07-ev-00479 James Napier v. Merck Case No.:06-cv-9451 Wilma Parslow v. Merck Case No.: 07-cv-3465 Tamae Parter v. Merck Case No.: 07-cv-790 Marguret Sizelove v. Merck Case No.: 07-cv-3459 Maxine Sklute v. Merck Case No.: 07-cv-3461 Edna Sykes v. Merck Case No.:07-cv-1412 Kathleen Toelke v. Merck Case No.: 06-cv-7629 Reyna Vandercar v. Merck Case No.:06-cv-11329 Nellie Wade v. Merck Case No.: 06-cv-9453 Carol Young v. Merck Case No.: 06-cy-0531 Bonnie Laughlin v. Merck Case No.:07-cv-3827



1:06-md-1789 (JFK)

ORDER FOR ADMISSION PRO HAC VICE

USDC SDNY
DOCUMENT
ELECTRONICALLY FILED
DOC #:
DATE FILED: 5-30-2007

Considering the Certificate of Good Standing submitted by counsel, IT IS

HEREBY ORDERED that:

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Meghan M. Tans
Levin, Papantonio, Thomas, Mitchell, Echsner & Proctor, P.A.
316 S. Baylen Street, Suite 400
P.O. Box 12308 (32591)
Pensacola, FL 32502
Phone: (850) 435-7181

Phone: (850) 435-7181
Facsimile: (850) 436-6181
Email: mtans@levinlaw.com

is admitted to practice pro hac vice as counsel for the plaintiffs in the above-referenced

MDL proceeding.

Signed this 30 day of

_, 2007, at

, New

York,

UNITED STATES DISTRICT COURT JUDGE

غمدومو بديد

Case 1:06-md-01789-JFK-JCF Document 110 Filed 05/30/2007 Page 3 of 3

VO 136 (Rev. 9 98) Centificate of Good Standing

UNITED STATES DISTRICT COURT

MIDDLE DISTRICT OF FLORIDA

CERTIFICATE OF GOOD STANDING

I, Sheryl L. Loesch, Clerk of this Court,
certify that Meghan M. Tans, Bar # 00888745,
was duly admitted to practice in this Court on
April 22, 2005, and is in good standing

Dated at Jacksonville, Florida, on May 17, 2007.

as a member of the Bur of this Court.

SHERYL L. LOESCII

CLERK

DEPUTY CLERK